California Drug Recall Information

Recall Name
Hospira, Inc. Recalls Three Lots of Propofol Injectable Due to Suspected Particles Embedded in Glass

Recall Date	Product Description	Recalling Firm	Recall Reason
8/14/12	Propofol Injectable Emulsion, 1%, 1g/100ml	**Hospira, Inc.** Lake Forest, IL	*Embedded particles may dislodge in solution*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Propofol Injectable Emulsion, 1%, NDC # 0409-4699-24 Suspect Lots Recalled : • 07-893-DJ, Exp. 01JUL2013 • 10-123-DJ, Exp. 01OCT2013 • 10-125-DJ, Exp. 01OCT2013	**CA**, nationwide	From September 2011 through February 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm315719.htm